-continued

Val	Val	35 35	Val	Ser	His	Glu	Asp 40	Pro	Glu	Val	Lys	Phe 45	Asn	Trp	Tyr
Val	Asp 50	Gly	Val	Glu	Val	His 55	Asn	Ala	Lys	Thr	60 Lys	Pro	Arg	Glu	Glu
Glr 65	Tyr	Asn	Ser	Thr	Tyr 70	Arg	Val	Val	Ser	Val 75	Leu	Thr	Val	Leu	His 80
Gln	Asp	Trp	Leu	Asn 85	Gly	Lys	Glu	Tyr	Lys 90	Cys	Lys	Val	Ser	Asn 95	Lys
Ala	Leu	Pro	Ala 100	Pro	Ile	Glu	Lys	Thr 105	Ile	Ser	ГÀа	Ala	Lys 110	Gly	Gln
Pro	Arg	Glu 115	Pro	Gln	Val	Tyr	Thr 120	Leu	Pro	Pro	Ser	Arg 125	Asp	Glu	Leu
Thr	Lys 130	Asn	Gln	Val	Ser	Leu 135	Thr	Cys	Leu	Val	Lys 140	Gly	Phe	Tyr	Pro
Ser 145	Asp	Ile	Ala	Val	Glu 150	Trp	Glu	Ser	Asn	Gly 155	Gln	Pro	Glu	Asn	Asn 160
Tyr	Lys	Thr	Thr	Pro 165	Pro	Val	Leu	Asp	Ser 170	Asp	Gly	Ser	Phe	Phe 175	Leu
Tyr	Ser	Lys	Leu 180	Thr	Val	Asp	Lys	Ser 185	Arg	Trp	Gln	Gln	Gly 190	Asn	Val
Phe	Ser	Сув 195	Ser	Val	Met	His	Glu 200	Ala	Leu	His	Asn	His 205	Tyr	Thr	Gln
Lys	Ser 210	Leu	Ser	Leu	Ser	Pro 215	Gly	Lys							

1-46. (canceled)

- 47. A method of treating a subject having an epidermal growth factor 2+ (HER2+) tumor that expresses HER2 at a 2+ level or lower as determined by immunohistochemistry (IHC), the method comprising administering to the subject an effective amount of a combination of a first and a second monovalent antigen-binding construct,
 - a) wherein the first and second monovalent antigenbinding constructs are distinct and each comprises a single antigen-binding polypeptide construct and a dimeric Fc comprising a first Fc polypeptide and a second Fc polypeptide each comprising a CH2 sequence and a CH3 sequence, the dimeric Fc coupled, with or without a linker, to the antigen-binding polypeptide construct, and wherein the first monovalent antigen-binding construct and the second monovalent antigen-binding construct is each selected from the group consisting of constructs 1 through 5, wherein:
 - i) construct 1 comprises an H-CDR1 comprising the sequence set forth in SEQ ID NO:179, an H-CDR2 comprising the sequence set forth in SEQ ID NO:183, an H-CDR3 comprising the sequence set forth in SEQ ID NO:181, an L-CDR1 comprising the sequence set forth in SEQ ID NO:245, an L-CDR2 comprising the sequence set forth in SEQ ID NO:249, and an L-CDR3 comprising the sequence set forth in SEQ ID NO:247, and
 - ii) construct 2 comprises an H-CDR1 comprising the sequence set forth in SEQ ID NO:195, an H-CDR2 comprising the sequence set forth in SEQ ID NO:199, an H-CDR3 comprising the sequence set forth in SEQ

- ID NO:197, an L-CDR1 comprising the sequence set forth in SEQ ID NO:245, an L-CDR2 comprising the sequence set forth in SEQ ID NO:249, and an L-CDR3 comprising the sequence set forth in SEQ ID NO:247, and
- iii) construct 3 comprises an H-CDR1 comprising the sequence set forth in SEQ ID NO:273, an H-CDR2 comprising the sequence set forth in SEQ ID NO:277, an H-CDR3 comprising the sequence set forth in SEQ ID NO:275, an L-CDR1 comprising the sequence set forth in SEQ ID NO:89, an L-CDR2 comprising the sequence set forth in SEQ ID NO:93, and an L-CDR3 comprising the sequence set forth in SEQ ID NO:91, and
- iv) construct 4 comprises an H-CDR1 comprising the sequence set forth in SEQ ID NO:225, an H-CDR2 comprising the sequence set forth in SEQ ID NO:229, an H-CDR3 comprising the sequence set forth in SEQ ID NO:227, an L-CDR1 comprising the sequence set forth in SEQ ID NO:217, an L-CDR2 comprising the sequence set forth in SEQ ID NO:221, and an L-CDR3 comprising the sequence set forth in SEQ ID NO:219, and
- v) construct 5 comprises an H-CDR1 comprising the sequence set forth in SEQ ID NO:101, an H-CDR2 comprising the sequence set forth in SEQ ID NO:105, an H-CDR3 comprising the sequence set forth in SEQ ID NO:103, an L-CDR1 comprising the sequence set forth in SEQ ID NO:109, an L-CDR2 comprising the